

09/778,926.

FILE 'HOME' ENTERED AT 15:49:47 ON 19 MAY 2005

=> file biosis medline caplus wpids uspatfull

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FILE 'BIOSIS' ENTERED AT 15:51:13 ON 19 MAY 2005

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FILE 'CAPLUS' ENTERED AT 15:51:13 ON 19 MAY 2005

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FILE 'USPATFULL' ENTERED AT 15:51:13 ON 19 MAY 2005

CA INDEXING COPYRIGHT (C) 2005 AMERICAN CHEMICAL SOCIETY (ACS)

\*\*\* YOU HAVE NEW MAIL \*\*\*

=> s bovine spongiform encephalopathy and electrophoresis

L1 554 BOVINE SPONGIFORM ENCEPHALOPATHY AND ELECTROPHORESIS

=> s l1 and glycoform

L2 9 L1 AND GLYCOFORM

=> dup rem l2

PROCESSING COMPLETED FOR L2

L3 9 DUP REM L2 (0 DUPLICATES REMOVED)

=> d l3 bib abs 1-9

L3 ANSWER 1 OF 9 USPATFULL on STN

AN 2004:334822 USPATFULL

TI Diagnostic method

IN Stack, Michael James, Surrey, UNITED KINGDOM

Chaplin, Melanie Jane, Surrey, UNITED KINGDOM

Clark, Jemma, Surrey, UNITED KINGDOM

PI US 2004265904 A1 20041230

AI US 2004-493572 A1 20040513 (10)

WO 2002-GB4789 20021023

PRAI GB 2001-25606 20011025

DT Utility

FS APPLICATION

LREP NIXON & VANDERHYE, PC, 1100 N GLEBE ROAD, 8TH FLOOR, ARLINGTON, VA,  
22201-4714

CLMN Number of Claims: 16

ECL Exemplary Claim: 1

DRWN 4 Drawing Page(s)

LN.CNT 692

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB A method for typing a strain of a transmissible spongiform encephalopathy (TSE) in an infected animal, said method comprising: a) separating a sample of abnormal prion protein on the basis of molecular weight and/or **glycoform** ratios, and detecting the separated forms; b) detecting in the sample the presence of a peptide sequence, wherein the presence of said peptide sequence within abnormal prion protein is capable of distinguishing a particular strain of TSE from others, and c) using the results of (a) and (b) to determine the type of TSE strain present in the sample. The method may be used in particular to distinguish BSE from scrapie in sheep.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L3 ANSWER 2 OF 9 USPATFULL on STN  
AN 2004:307835 USPATFULL  
TI Method  
IN Fisher, Elizabeth Mary Claire, London, UNITED KINGDOM  
Lloyd, Sarah Elizabeth, London, UNITED KINGDOM  
Collinge, John, Queen Square, UNITED KINGDOM  
PI US 2004242511 A1 20041202  
AI US 2004-470014 A1 20040122 (10)  
WO 2002-GB256 20020122  
PRAI GB 2001-1763 20010123  
DT Utility  
FS APPLICATION  
LREP MARSHALL, GERSTEIN & BORUN LLP, 6300 SEARS TOWER, 233 S. WACKER DRIVE,  
CHICAGO, IL, 60606  
CLMN Number of Claims: 31  
ECL Exemplary Claim: 1  
DRWN No Drawings  
LN.CNT 3578

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The invention relates to a method for the detection of prions in a sample comprising the steps of contacting one or more test animals with the sample; incubating the test animals; monitoring the test animals for adverse effects or death; and optionally performing a biopsy on the test animals that display adverse effects or death for evidence of prions; wherein the test animals have prion incubation times of 196 days or less.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L3 ANSWER 3 OF 9 USPATFULL on STN  
AN 2004:222053 USPATFULL  
TI Endomannosidases in the modification of glycoproteins in eukaryotes  
IN Hamilton, Stephen, Enfield, NH, UNITED STATES  
PI US 2004171826 A1 20040902  
AI US 2003-695243 A1 20031027 (10)  
RLI Continuation-in-part of Ser. No. US 2003-371877, filed on 20 Feb 2003,  
PENDING  
DT Utility  
FS APPLICATION  
LREP James F. Haley, Jr., Esq., c/o FISH & NEAVE, 1251 Avenue of the  
Americas, New York, NY, 10020-1104  
CLMN Number of Claims: 25  
ECL Exemplary Claim: 1  
DRWN 15 Drawing Page(s)  
LN.CNT 2983

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The present invention generally relates to methods of modifying the glycosylation structures of recombinant proteins expressed in fungi or other lower eukaryotes, to more closely resemble the glycosylation of proteins from higher mammals, in particular humans. The present invention also relates to novel enzymes and, nucleic acids encoding them and, hosts engineered to express the enzymes, methods for producing modified glycoproteins in hosts and modified glycoproteins so produced.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L3 ANSWER 4 OF 9 USPATFULL on STN  
AN 2004:171948 USPATFULL  
TI Method  
IN Enari, Masato, Chuo-ku, JAPAN  
Flechsigs, Eckhard, Versbacher, GERMANY, FEDERAL REPUBLIC OF  
Collinge, John, Queen, UNITED KINGDOM  
Weismann, Charles, London, UNITED KINGDOM  
PI US 2004132109 A1 20040708  
AI US 2004-470022 A1 20040109 (10)  
WO 2002-GB257 20020122

PRAI GB 2001-1762 20010123  
DT Utility  
FS APPLICATION  
LREP MARSHALL, GERSTEIN & BORUN LLP, 6300 SEARS TOWER, 233 S. WACKER DRIVE,  
CHICAGO, IL, 60606  
CLMN Number of Claims: 31  
ECL Exemplary Claim: 1  
DRWN No Drawings  
LN.CNT 3141

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The invention relates to methods for determining the presence of prions in a tissue/organ or fluid therefrom; said method comprising the steps of: contacting the tissue/organ with one or more devices, wherein said devices are capable of binding prions; removing said devices from contact with said tissue/organ; determining if said devices are binding prions wherein the device is contacted with the tissue/organ for 120 minutes.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L3 ANSWER 5 OF 9 USPATFULL on STN  
AN 2003:60077 USPATFULL  
TI Immunological agents specific for prion protein (PRP)  
IN Sy, Man-Sun, Shaker Heights, OH, United States  
Gambetti, Pierluigi, Shaker Heights, OH, United States  
PA Case Western Reserve University, Cleveland, OH, United States (U.S. corporation)  
PI US 6528269 B1 20030304  
AI US 1998-204816 19981203 (9)  
PRAI US 1998-90165P 19980622 (60)  
DT Utility  
FS GRANTED  
EXNAM Primary Examiner: Swartz, Rodney P  
LREP Fay, Sharpe, Fagan, Minnich & McKee, LLP  
CLMN Number of Claims: 4  
ECL Exemplary Claim: 1  
DRWN 15 Drawing Figure(s); 15 Drawing Page(s)  
LN.CNT 1433

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The present invention is directed to a panel of monoclonal antibodies (Mabs) specific for murine prion protein PrP.sup.c. These Mabs can be applied to immunoblotting, cell surface immunofluorescent staining and immunohistochemistry at light and electron microscopy. Additionally, these Mabs recognize both the normal (PrP.sup.c) and protease-resistant (PrP.sup.res) isoforms of PrP. Some Mabs are species restricted, while others react with PrP from a broad range of mammals including mice, humans, monkeys, cows, sheep, squirrels and hamsters. Moreover, several of the Mabs selectively recognize different PrP glycoforms as well as the metabolic fragments of PrP.sup.c. These newly generated PrP.sup.c antibodies are useful for exploring the biology of PrP.sup.c and to establish the diagnosis of prion diseases in both humans and animals.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L3 ANSWER 6 OF 9 USPATFULL on STN  
AN 2002:251169 USPATFULL  
TI Method of detecting PrP protein and kits therefor  
IN Voelkel, Dirk, Vienna, AUSTRIA  
Zimmermann, Klaus, Vienna, AUSTRIA  
Turecek, Peter, Klosterneuburg, AUSTRIA  
Schwarz, Hans-Peter, Vienna, AUSTRIA  
PI US 2002137114 A1 20020926  
AI US 2002-51413 A1 20020118 (10)  
PRAI US 2001-263022P 20010119 (60)  
DT Utility  
FS APPLICATION  
LREP Baxter Healthcare Corporation, P.O. Box 15210, Irvine, CA, 92614  
CLMN Number of Claims: 49

ECL Exemplary Claim: 1

DRWN 7 Drawing Page(s)

LN.CNT 997

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The invention relates to a method for the detection of neurological disorders in a patient comprising (a) measuring the concentration of PrP protein in a biological fluid sample of said patient; and (b) determining whether said concentration of said PrP protein is above or below a predetermined threshold value, whereby the concentration above said predetermined threshold value identifies a patient with a neurological disorder, a method for the detection and quantification of PrP protein and pathogenic PrP<sup>sup.res</sup> protein in a sample, and a kit comprising a set of reagents to determine the concentration of PrP protein and pathogenic PrP<sup>sup.res</sup> protein in a sample.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L3 ANSWER 7 OF 9 USPATFULL on STN

AN 2002:157046 USPATFULL

TI Diagnosis of spongiform encephalopathy

IN Collinge, John, London, UNITED KINGDOM

PI US 2002081645 A1 20020627

AI US 2001-778926 A1 20010206 (9)

RLI Continuation of Ser. No. US 1999-291215, filed on 14 Apr 1999, ABANDONED

PRAI GB 1996-21469 19961015

GB 1996-21885 19961021

DT Utility

FS APPLICATION

LREP HALE AND DORR, LLP, 60 STATE STREET, BOSTON, MA, 02109

CLMN Number of Claims: 34

ECL Exemplary Claim: 1

DRWN 9 Drawing Page(s)

LN.CNT 1149

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The present invention relates to a method for typing a sample of a prion or spongiform encephalopathy disease, a kit suitable for use in such a typing method, a method for identifying infection in an animal and/or tissue of **bovine spongiform encephalopathy** (BSE), a method for assessing and/or predicting the susceptibility of an animal to BSE, a kit for use in such an assessment and/or prediction method, a method for the treatment of a prion disease, and compounds suitable for such a method.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L3 ANSWER 8 OF 9 USPATFULL on STN

AN 2002:78467 USPATFULL

TI Mammalian proteins; related reagents and methods

IN Bazan, J. Fernando, Palo Alto, CA, UNITED STATES

PI US 2002042122 A1 20020411

AI US 2000-745003 A1 20001220 (9)

PRAI US 1999-172090P 19991223 (60)

DT Utility

FS APPLICATION

LREP DNAX RESEARCH INSTITUTE, LEGAL DEPARTMENT, 901 CALIFORNIA AVENUE, PALO ALTO, CA, 94304

CLMN Number of Claims: 20

ECL Exemplary Claim: 1

DRWN No Drawings

LN.CNT 2359

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB Mammalian polypeptides, isolated proteins, and fragments thereof including the polynucleotides encoding them. Antibodies, both polyclonal and monoclonal, are also provided. Methods of using the compositions for both diagnostic and therapeutic utilities are provided.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L3 ANSWER 9 OF 9 MEDLINE on STN  
 AN 2000237810 MEDLINE  
 DN PubMed ID: 10773427.  
 TI Comparison of French natural scrapie isolates with **bovine spongiform encephalopathy** and experimental scrapie infected sheep.  
 AU Baron T G; Madec J Y; Calavas D; Richard Y; Barillet F  
 CS Agence Francaise de Securite Sanitaire des Aliments, 69364, Lyon, France..  
 t.baron@lyon.afssa.fr  
 SO Neuroscience letters, (2000 Apr 28) 284 (3) 175-8.  
 Journal code: 7600130. ISSN: 0304-3940.  
 CY Ireland  
 DT Journal; Article; (JOURNAL ARTICLE)  
 LA English  
 FS Priority Journals  
 EM 200006  
 ED Entered STN: 20000622  
 Last Updated on STN: 20000622  
 Entered Medline: 20000613  
 AB We compared the **glycoform** pattern of the abnormal prion protein (PrP(Sc)) detected by immunoblotting in 21 sheep with natural scrapie, from 21 different outbreaks identified in France since 1996, with a **bovine spongiform encephalopathy** (BSE)-infected sheep. All the natural scrapie isolates had a higher molecular mass of the unglycosylated PrP(Sc) than in BSE-infected sheep. In the latter case, this molecular mass appeared identical to that found in the CH 1641 experimental scrapie strain (type C pattern), whereas in natural scrapie cases it was similar to that found in the SSBP/1 experimental scrapie strains. These results suggest that all French natural scrapie isolates studied so far would belong, as SSBP/1, to the group of scrapie cases with type A electrophoretic pattern.

=> d his

(FILE 'HOME' ENTERED AT 15:49:47 ON 19 MAY 2005)

FILE 'BIOSIS, MEDLINE, CAPLUS, WPIDS, USPATFULL' ENTERED AT 15:51:13 ON 19 MAY 2005

L1 554 S BOVINE SPONGIFORM ENCEPHALOPATHY AND ELECTROPHORESIS  
 L2 9 S L1 AND GLYCOFORM  
 L3 9 DUP REM L2 (0 DUPLICATES REMOVED)

=> s prion and typing'

MISMATCHED QUOTE 'TYPING''

Quotation marks (or apostrophes) must be used in pairs, one before and one after the expression you are setting off or masking.

=> s prion and typing

L4 479 PRION AND TYPING

=> s l4 and glycoform?

L5 21 L4 AND GLYCOFORM?

=> s l5 and electrophoresis

L6 15 L5 AND ELECTROPHORESIS

=> dup rem 16

PROCESSING COMPLETED FOR L6

L7 15 DUP REM L6 (0 DUPLICATES REMOVED)

=> s l7 and ratio?

L8 13 L7 AND RATIO?

=> d l8 bib abs 1-13

L8 ANSWER 1 OF 13 CAPLUS COPYRIGHT 2005 ACS on STN

AN 2003:335409 CAPLUS  
 DN 138:317152  
 TI Diagnostic method  
 IN Stack, Michael James; Chaplin, Melanie Jane; Clark, Jemma  
 PA The Secretary of State for Environment, Food and Rural Affairs, UK  
 SO PCT Int. Appl., 30 pp.  
 CODEN: PIXXD2  
 DT Patent  
 LA English  
 FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2003036303	A1	20030501	WO 2002-GB4789	20021023
	WO 2003036303	C1	20030918		
	W:	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW			
	RW:	GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG			
	CA 2462581	AA	20030501	CA 2002-2462581	20021023
	GB 2396009	A1	20040609	GB 2004-6547	20021023
	GB 2396009	B2	20050316		
	EP 1442303	A1	20040804	EP 2002-770097	20021023
	R:	AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, SK			
	JP 2005506551	T2	20050303	JP 2003-538748	20021023
	US 2004265904	A1	20041230	US 2004-493572	20040513
PRAI	GB 2001-25606	A	20011025		
	WO 2002-GB4789	W	20021023		

AB A method for **typing** a strain of a transmissible spongiform encephalopathy (TSE) in an infected animal, said method comprising: (a) separating a sample of abnormal **prion** protein on the basis of mol. weight and/or **glycoform ratios**, and detecting the separated forms; (b) detecting in the sample the presence of a peptide sequence, wherein the presence of said peptide sequence within abnormal **prion** protein is capable of distinguishing a particular strain of TSE from others, and (c) using the results of (a) and (b) to determine the type of TSE strain present in the sample. The method may be used in particular to distinguish BSE from scrapie in sheep.

RE.CNT 8 THERE ARE 8 CITED REFERENCES AVAILABLE FOR THIS RECORD  
 ALL CITATIONS AVAILABLE IN THE RE FORMAT

L8 ANSWER 2 OF 13 USPATFULL on STN  
 AN 2005:107273 USPATFULL  
 TI Method to type **prion** proteins  
 IN Collinge, John, London, UNITED KINGDOM  
 Wadsworth, Jonathan David Frank, London, UNITED KINGDOM  
 PA D-Gen Limited, London, UNITED KINGDOM (non-U.S. corporation)  
 PI US 6887676 B1 20050503  
 WO 2000062068 20001019  
 AI US 2001-958517 20000407 (9)  
 WO 2000-GB1327 20000407  
 20020212 PCT 371 date  
 PRAI GB 2001-9908059 19990409  
 DT Utility  
 FS GRANTED  
 EXNAM Primary Examiner: Housel, James; Assistant Examiner: Lucas, Zachariah  
 LREP Nikolai & Mersereau PA, Mersereau, C. G.  
 CLMN Number of Claims: 17  
 ECL Exemplary Claim: 1  
 DRWN 4 Drawing Figure(s); 4 Drawing Page(s)  
 LN.CNT 983  
 AB The invention relates to methods and materials for use in the

**typing**, diagnosis, prevention and/or treatment of **prion** disease.

L8 ANSWER 3 OF 13 USPATFULL on STN  
AN 2005:63014 USPATFULL  
TI Albumin fusion proteins  
IN Rosen, Craig A., Laytonsville, MD, UNITED STATES  
Haseltine, William A., Washington, DC, UNITED STATES  
PA Human Genome Sciences, Inc. (U.S. corporation)  
PI US 2005054051 A1 20050310  
AI US 2004-922142 A1 20040820 (10)  
RLI Division of Ser. No. US 2001-832929, filed on 12 Apr 2001, PENDING  
DT Utility  
FS APPLICATION  
LREP FINNEGAN, HENDERSON, FARABOW, GARRETT & DUNNER, LLP, 1300 I STREET, NW,  
WASHINGTON, DC, 20005  
CLMN Number of Claims: 33  
ECL Exemplary Claim: 1  
DRWN 20 Drawing Page(s)  
LN.CNT 17526

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The present invention encompasses albumin fusion proteins. Nucleic acid molecules encoding the albumin fusion proteins of the invention are also encompassed by the invention, as are vectors containing these nucleic acids, host cells transformed with these nucleic acids vectors, and methods of making the albumin fusion proteins of the invention and using these nucleic acids, vectors, and/or host cells. Additionally the present invention encompasses pharmaceutical compositions comprising albumin fusion proteins and methods of treating, preventing, or ameliorating diseases, disorders or conditions using albumin fusion proteins of the invention.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L8 ANSWER 4 OF 13 USPATFULL on STN  
AN 2004:334822 USPATFULL  
TI Diagnostic method  
IN Stack, Michael James, Surrey, UNITED KINGDOM  
Chaplin, Melanie Jane, Surrey, UNITED KINGDOM  
Clark, Jemma, Surrey, UNITED KINGDOM  
PI US 2004265904 A1 20041230  
AI US 2004-493572 A1 20040513 (10)  
WO 2002-GB4789 20021023  
PRAI GB 2001-25606 20011025  
DT Utility  
FS APPLICATION  
LREP NIXON & VANDERHYE, PC, 1100 N GLEBE ROAD, 8TH FLOOR, ARLINGTON, VA,  
22201-4714  
CLMN Number of Claims: 16  
ECL Exemplary Claim: 1  
DRWN 4 Drawing Page(s)  
LN.CNT 692

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB A method for **typing** a strain of a transmissible spongiform encephalopathy (TSE) in an infected animal, said method comprising: a) separating a sample of abnormal **prion** protein on the basis of molecular weight and/or **glycoform ratios**, and detecting the separated forms; b) detecting in the sample the presence of a peptide sequence, wherein the presence of said peptide sequence within abnormal **prion** protein is capable of distinguishing a particular strain of TSE from others, and c) using the results of (a) and (b) to determine the type of TSE strain present in the sample. The method may be used in particular to distinguish BSE from scrapie in sheep.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L8 ANSWER 5 OF 13 USPATFULL on STN  
AN 2004:314570 USPATFULL  
TI 47153, A HUMAN GLYCOSYLTRANSFERASE FAMILY MEMBER AND USES THEREFOR  
IN Meyers, Rachel, Newton, MA, UNITED STATES  
Rosenfeld, Julie Beth, Sharon, MA, UNITED STATES  
PI US 2004248242 A1 20041209  
US 6849437 B2 20050201  
AI US 2002-113709 A1 20020328 (10)  
PRAI US 2001-279647P 20010328 (60)  
DT Utility  
FS APPLICATION  
LREP Intellectual Property Group, MILLENIUM PHARMACEUTICALS, INC., 75 Sidney  
Street, Cambridge, MA, 02139  
CLMN Number of Claims: 18  
ECL Exemplary Claim: 1  
DRWN 8 Drawing Page(s)  
LN.CNT 4650

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The invention provides isolated nucleic acids molecules, designated  
47153 nucleic acid molecules, which encode novel glycosyltransferase  
family members. The invention also provides antisense nucleic acid  
molecules, recombinant expression vectors containing 47153 nucleic acid  
molecules, host cells into which the expression vectors have been  
introduced, and nonhuman transgenic animals in which a 47153 gene has  
been introduced or disrupted. The invention still further provides  
isolated 47153 proteins, fusion proteins, antigenic peptides and  
anti-47153 antibodies. Diagnostic and therapeutic methods utilizing  
compositions of the invention are also provided.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L8 ANSWER 6 OF 13 USPATFULL on STN  
AN 2004:221354 USPATFULL  
TI ALBUMIN FUSION PROTEINS  
IN Rosen, Craig A., Laytonsville, MD, UNITED STATES  
Haseltine, William A., Washington, DC, UNITED STATES  
PI US 2004171123 A1 20040902  
AI US 2001-832929 A1 20010412 (9)  
DT Utility  
FS APPLICATION  
LREP FINNEGAN, HENDERSON, FARABOW, GARRETT & DUNNER, LLP, 1300 I STREET, NW,  
WASHINGTON, DC, 20005  
CLMN Number of Claims: 29  
ECL Exemplary Claim: 1  
DRWN 18 Drawing Page(s)  
LN.CNT 17424

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The present invention encompasses albumin fusion proteins. Nucleic acid  
molecules encoding the albumin fusion proteins of the invention are also  
encompassed by the invention, as are vectors containing these nucleic  
acids, host cells transformed with these nucleic acids vectors, and  
methods of making the albumin fusion proteins of the invention and using  
these nucleic acids, vectors, and/or host cells. Additionally the  
present invention encompasses pharmaceutical compositions comprising  
albumin fusion proteins and methods of treating, preventing, or  
ameliorating diseases, disorders or conditions using albumin fusion  
proteins of the invention.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L8 ANSWER 7 OF 13 USPATFULL on STN  
AN 2004:107607 USPATFULL  
TI 47174, a novel human glycosyltransferase and uses thereof  
IN Kapeller-Libermann, Rosana, Chestnut Hill, MA, UNITED STATES  
PA Millennium Pharmaceuticals, Inc. (U.S. corporation)  
PI US 2004082007 A1 20040429  
AI US 2003-713345 A1 20031114 (10)  
RLI Division of Ser. No. US 2001-973457, filed on 9 Oct 2001, GRANTED, Pat.



No. US 6703230  
PRAI US 2000-238849P 20001006 (60)  
DT Utility  
FS APPLICATION  
LREP MILLENNIUM PHARMACEUTICALS, INC., 75 Sidney Street, Cambridge, MA, 02139  
CLMN Number of Claims: 20  
ECL Exemplary Claim: 1  
DRWN 4 Drawing Page(s)  
LN.CNT 4889

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The invention provides isolated nucleic acids molecules, designated 47174 nucleic acid molecules, which encode novel glycosyltransferase family members. The invention also provides antisense nucleic acid molecules, recombinant expression vectors containing 47174 nucleic acid molecules, host cells into which the expression vectors have been introduced, and nonhuman transgenic animals in which a 47174 gene has been introduced or disrupted. The invention still further provides isolated 47174 proteins, fusion proteins, antigenic peptides and anti-47174 antibodies. Diagnostic methods utilizing compositions of the invention are also provided. The invention also provides methods of modulating pain or pain related disorders utilizing the compositions of the invention. Accordingly, methods of treating, preventing and/or diagnosing neurological disorders are disclosed.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L8 ANSWER 8 OF 13 USPATFULL on STN  
AN 2003:318632 USPATFULL  
TI Novel human transferase family members and uses thereof  
IN Meyers, Rachel E., Newton, MA, UNITED STATES  
Williamson, Mark, Saugus, MA, UNITED STATES  
Leiby, Kevin R., Natick, MA, UNITED STATES  
Kapeller-Libermann, Rosana, Chestnut Hill, MA, UNITED STATES  
Olandt, Peter J., Newton, MA, UNITED STATES  
MacBeth, Kyle J., Boston, MA, UNITED STATES  
Rudolph-Owen, Laura A., Jamaica Plain, MA, UNITED STATES  
Tsai, Fong-Ying, Newton, MA, UNITED STATES  
Hunter, John J., Somerville, MA, UNITED STATES

PI US 2003224376 A1 20031204

AI US 2002-184648 A1 20020627 (10)

RLI Continuation-in-part of Ser. No. US 2001-815028, filed on 22 Mar 2001, PENDING Continuation-in-part of Ser. No. US 2001-801220, filed on 7 Mar 2001, PENDING Continuation-in-part of Ser. No. US 2001-816714, filed on 23 Mar 2001, ABANDONED Continuation-in-part of Ser. No. US 2001-844948, filed on 27 Apr 2001, PENDING Continuation-in-part of Ser. No. US 2001-861164, filed on 18 May 2001, ABANDONED Continuation-in-part of Ser. No. US 2001-883060, filed on 15 Jun 2001, PENDING Continuation-in-part of Ser. No. US 2001-962678, filed on 25 Sep 2001, PENDING Continuation-in-part of Ser. No. US 2001-973457, filed on 9 Oct 2001, PENDING Continuation-in-part of Ser. No. US 2002-72285, filed on 8 Feb 2002, PENDING Continuation-in-part of Ser. No. US 2001-817910, filed on 26 Mar 2001, PENDING Continuation-in-part of Ser. No. US 2001-842528, filed on 25 Apr 2001, ABANDONED Continuation-in-part of Ser. No. US 2001-882836, filed on 15 Jun 2001, PENDING Continuation-in-part of Ser. No. US 2001-882872, filed on 15 Jun 2001, ABANDONED

PRAI WO 2001-US9358 20010322  
WO 2001-US7269 20010307  
WO 2001-US9468 20010323  
WO 2001-US13805 20010427  
WO 2001-US16292 20010518  
WO 2001-US19138 20010615  
WO 2001-US29963 20010925  
WO 2002-US3736 20020208  
WO 2001-US9633 20010326  
WO 2001-US40607 20010425  
WO 2001-US19543 20010615  
WO 2001-US19153 20010615  
US 2000-191964P 20000324 (60)

US 2000-187456P 20000307 (60)  
US 2000-191865P 20000324 (60)  
US 2000-200604P 20000428 (60)  
US 2000-205408P 20000519 (60)  
US 2000-212079P 20000615 (60)  
US 2000-235044P 20000925 (60)  
US 2000-238849P 20001006 (60)  
US 2001-267494P 20010208 (60)  
US 2000-192092P 20000324 (60)  
US 2000-199500P 20000425 (60)  
US 2000-211730P 20000615 (60)  
US 2000-212077P 20000615 (60)

DT Utility

FS APPLICATION

LREP Theodore R. Allen, Millennium Pharmaceuticals, Inc., 75 Sidney Street,  
Cambridge, MA, 02139

CLMN Number of Claims: 19

ECL Exemplary Claim: 1

DRWN 125 Drawing Page(s)

LN.CNT 66695

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The invention provides isolated nucleic acids molecules, designated 33877, 47179, 26886, 25552, 32132, 32244, 23680, 32624, 47174, 60491, 46743, 27417, 27960, 32252, and 53320 nucleic acid molecules, which encode novel human transferase family members. The invention also provides antisense nucleic acid molecules, recombinant expression vectors containing 33877, 47179, 26886, 25552, 32132, 32244, 23680, 32624, 47174, 60491, 46743, 27417, 27960, 32252, or 53320 nucleic acid molecules, host cells into which the expression vectors have been introduced, and nonhuman transgenic animals in which a 33877, 47179, 26886, 25552, 32132, 32244, 23680, 32624, 47174, 60491, 46743, 27417, 27960, 32252, or 53320 gene has been introduced or disrupted. The invention still further provides isolated 33877, 47179, 26886, 25552, 32132, 32244, 23680, 32624, 47174, 60491, 46743, 27417, 27960, 32252, or 53320 proteins, fusion proteins, antigenic peptides and anti-33877, 47179, 26886, 25552, 32132, 32244, 23680, 32624, 47174, 60491, 46743, 27417, 27960, 32252, or 53320 antibodies. Diagnostic methods utilizing compositions of the invention are also provided.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L8 ANSWER 9 OF 13 USPATFULL on STN

AN 2002:294703 USPATFULL

TI 47174, a novel human glycosyltransferase and uses thereof

IN Kapeller-Libermann, Rosana, Chestnut Hill, MA, UNITED STATES

PI US 2002164746 A1 20021107

US 6703230 B2 20040309

AI US 2001-973457 A1 20011009 (9)

PRAI US 2000-238849P 20001006 (60)

DT Utility

FS APPLICATION

LREP LOUIS MEYERS, FISH & RICHARDSON P.C., 225 Franklin Street, Boston, MA,  
02110-2804

CLMN Number of Claims: 20

ECL Exemplary Claim: 1

DRWN 4 Drawing Page(s)

LN.CNT 4577

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The invention provides isolated nucleic acids molecules, designated 47174 nucleic acid molecules, which encode novel glycosyltransferase family members. The invention also provides antisense nucleic acid molecules, recombinant expression vectors containing 47174 nucleic acid molecules, host cells into which the expression vectors have been introduced, and nonhuman transgenic animals in which a 47174 gene has been introduced or disrupted. The invention still further provides isolated 47174 proteins, fusion proteins, antigenic peptides and anti-47174 antibodies. Diagnostic methods utilizing compositions of the invention are also provided. The invention also provides methods of

modulating pain or pain related disorders utilizing the compositions of the invention. Accordingly, methods of treating, preventing and/or diagnosing neurological disorders are disclosed.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L8 ANSWER 10 OF 13 USPATFULL on STN  
AN 2002:280065 USPATFULL  
TI 32624, a novel human UDP-glucuronosyl and glycosyl transferase family member and uses thereof  
IN Leiby, Kevin R., Natick, MA, UNITED STATES  
PI US 2002155499 A1 20021024  
AI US 2001-962678 A1 20010925 (9)  
PRAI US 2000-235044P 20000925 (60)  
DT Utility  
FS APPLICATION  
LREP LOUIS MYERS, FISH & RICHARDSON P.C., 225 Franklin Street, Boston, MA, 02110-2804  
CLMN Number of Claims: 20  
ECL Exemplary Claim: 1  
DRWN 3 Drawing Page(s)  
LN.CNT 5149

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The invention provides isolated nucleic acids molecules, designated 32624 nucleic acid molecules, which encode novel UDP-glucuronosyl and glycosyl transferase members. The invention also provides antisense nucleic acid molecules, recombinant expression vectors containing 32624 nucleic acid molecules, host cells into which the expression vectors have been introduced, and nonhuman transgenic animals in which a 32624 gene has been introduced or disrupted. The invention still further provides isolated 32624 proteins, fusion proteins, antigenic peptides and anti-32624 antibodies. Diagnostic methods utilizing compositions of the invention are also provided.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L8 ANSWER 11 OF 13 USPATFULL on STN  
AN 2002:228315 USPATFULL  
TI 32626, a novel human UDP-glycosyltransferase and uses thereof  
IN Leiby, Kevin R., Natick, MA, UNITED STATES  
Spaltmann, Frank, Cambridge, MA, UNITED STATES  
Cook, William James, Natick, MA, UNITED STATES  
PI US 2002123475 A1 20020905  
AI US 2001-895728 A1 20010629 (9)  
PRAI US 2000-215749P 20000630 (60)  
DT Utility  
FS APPLICATION  
LREP Carolyn A. Favorito, Morrison & Foerster LLP, Suite 500, 3811 Valley Centre Drive, San Diego, CA, 92130-2332  
CLMN Number of Claims: 23  
ECL Exemplary Claim: 1  
DRWN 5 Drawing Page(s)  
LN.CNT 4203

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The invention provides isolated nucleic acids molecules, designated 32626 nucleic acid molecules, which encode novel UDP-glycosyltransferase family members. The invention also provides antisense nucleic acid molecules, recombinant expression vectors containing 32626 nucleic acid molecules, host cells into which the expression vectors have been introduced, and nonhuman transgenic animals in which a 32626 gene has been introduced or disrupted. The invention still further provides isolated 32626 proteins, fusion proteins, antigenic peptides and anti-32626 antibodies. Diagnostic methods utilizing compositions of the invention are also provided.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L8 ANSWER 12 OF 13 USPATFULL on STN

AN 2002:199265 USPATFULL  
TI 26199, 33530, 33949, 47148, 50226, and 58764, novel human transferase  
family members and uses therefor  
IN Meyers, Rachel, Newton, MA, UNITED STATES  
MacBeth, Kyle, Boston, MA, UNITED STATES  
PI US 2002107376 A1 20020808  
AI US 2001-924358 A1 20010806 (9)  
PRAI US 2000-229300P 20000901 (60)  
DT Utility  
FS APPLICATION  
LREP Intellectual Property Group, MILLENNIUM PHARMACEUTICALS, INC, 75 Sidney  
Street, Cambridge, MA, 02139  
CLMN Number of Claims: 24  
ECL Exemplary Claim: 1  
DRWN 28 Drawing Page(s)  
LN.CNT 6380

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The invention provides isolated nucleic acids molecules, designated  
26199, 33530, 33949, 47148, 50226, or 58764 nucleic acid molecules,  
which encode novel transferase family members. The invention also  
provides antisense nucleic acid molecules, recombinant expression  
vectors containing 26199, 33530, 33949, 47148, 50226, or 58764 nucleic  
acid molecules, host cells into which the expression vectors have been  
introduced, and nonhuman transgenic animals in which a 26199, 33530,  
33949, 47148, 50226, or 58764 gene has been introduced or disrupted. The  
invention still further provides isolated 26199, 33530, 33949, 47148,  
50226, or 58764 proteins, fusion proteins, antigenic peptides and  
anti-26199, -33530, -33949, -47148, -50226, or -58764 antibodies.  
Diagnostic methods utilizing compositions of the invention are also  
provided.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L8 ANSWER 13 OF 13 USPATFULL on STN  
AN 2002:157046 USPATFULL  
TI Diagnosis of spongiform encephalopathy  
IN Collinge, John, London, UNITED KINGDOM  
PI US 2002081645 A1 20020627  
AI US 2001-778926 A1 20010206 (9)  
RLI Continuation of Ser. No. US 1999-291215, filed on 14 Apr 1999, ABANDONED  
PRAI GB 1996-21469 19961015  
GB 1996-21885 19961021  
DT Utility  
FS APPLICATION  
LREP HALE AND DORR, LLP, 60 STATE STREET, BOSTON, MA, 02109  
CLMN Number of Claims: 34  
ECL Exemplary Claim: 1  
DRWN 9 Drawing Page(s)  
LN.CNT 1149

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The present invention relates to a method for **typing** a sample  
of a **prion** or spongiform encephalopathy disease, a kit  
suitable for use in such a **typing** method, a method for  
identifying infection in an animal and/or tissue of bovine spongiform  
encephalopathy (BSE), a method for assessing and/or predicting the  
susceptibility of an animal to BSE, a kit for use in such an assessment  
and/or prediction method, a method for the treatment of a **prion**  
disease, and compounds suitable for such a method.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

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